

Claims

1. A pharmaceutical formulation comprising a pharmaceutical acceptable salt of glycopyrronium, a solvates or physiologically functional derivative thereof in combination with ciclesonide, a pharmaceutically acceptable salt, solvates or physiologically functional derivative thereof and a pharmaceutically acceptable carrier and/or one or more excipients, and optionally one or more other therapeutic ingredients.
2. Formulation according to claim 1, wherein pharmaceutical acceptable salt of glycopyrronium and ciclesonide are contained in the same pharmaceutical formulation (fixed combination).
3. Formulation according to claim 1, wherein pharmaceutical acceptable salt of glycopyrronium and ciclesonide are contained in different pharmaceutical formulations (free combination).
4. Formulation according to claim 1, comprising a compound selected from the group of [11 β ,16 α (R)]-16,17-[(Cyclohexylmethylen)bis(oxy)]-11-hydroxy-21-(2-methyl-1-oxopropoxy)pregna-1,4-dien-3,20-dion, [11 β ,16 α (S)]-16,17-[(Cyclohexylmethylen)bis(oxy)]-11-hydroxy-21-(2-methyl-1-oxopropoxy)pregna-1,4-dien-3,20-dion, [11 β ,16 α (R,S)]-16,17-[(Cyclohexylmethylen)bis(oxy)]-11-hydroxy-21-(2-methyl-1-oxopropoxy)pregna-1,4-dien-3,20-dion, 16 α ,17-(22R)-Cyclohexylmethylenedioxy-11 β ,21-dihydroxypregna-1,4-dien-3,20-dion, 16 α ,17-(22S)-Cyclohexylmethylenedioxy-11 β ,21-dihydroxypregna-1,4-dien-3,20-dion and 16 α ,17-(22R,S)-Cyclohexylmethylenedioxy-11 β ,21-dihydroxypregna-1,4-dien-3,20-dion.
5. Formulation according to claim 1, wherein the pharmaceutical acceptable salt of glycopyrronium is selected from the group of compounds racemic forms [S,S-, S,R, R,S- and R,R-forms] of the pharmaceutical acceptable salt of glycopyrronium in any mixing ratio and enantiomerically enriched S,S-, S,R, R,S- and R,R-forms of the pharmaceutical acceptable salt of glycopyrronium.
6. Formulation according to claim 5, wherein the enantiomerically enriched form of the pharmaceutical acceptable salt of glycopyrronium is the R,R-form (i.e. (3R,2'R)-3-[(cyclopentylhydroxyphenylacetyl)oxy]-1,1-dimethylpyrrolidinium).
7. Formulation according to claim 6, wherein the R,R-form has an enantiomeric purity of 90% minimum enantiomeric excess (ee), preferably 95 % ee, more preferably more than 98 % ee, and in particular preferably more than 99.5 % ee.

8. Formulation according to claim 1 wherein the pharmaceutical acceptable salt of glycopyrronium is (3R,2'R)-3-[(cyclopentylhydroxyphenylacetyl)oxy]-1,1-dimethylpyrrolidinium bromide, which substantially does not contain glycopyrronium in the S,S-, S,R- and/or R,S- forms.
9. Formulation according to claim 1, comprising pharmaceutical acceptable salt of glycopyrronium and ciclesonide in an amount and ratio to be effective for a twice or once daily treatment of a clinical condition in a mammal, such as a human, for which a corticosteroid and/or an anticholinergic agent is indicated.
10. Formulation according to claim 1, which is suitable for administration by inhalation.
11. Formulation according to claim 1, which is suitable for nasal administration.
12. Pharmaceutical formulation according to claim 1, which is a dry powder and the carrier is a saccharide.
13. Pharmaceutical formulation according to claim 12, wherein the carrier is lactose monohydrate.
14. Method for the prophylaxis or treatment of a clinical condition in a mammal, such as a human, for which a corticosteroid and/or an anticholinergic agent is indicated, which comprises administration of a therapeutically effective amount of a pharmaceutical formulation comprising ciclesonide or a pharmaceutical acceptable salt, solvate, or physiologically functional derivative thereof in combination with a pharmaceutical acceptable salt of glycopyrronium, a solvate, or physiologically functional derivative thereof, and a pharmaceutical acceptable carrier and/or one or more excipients.
15. Method according to claim 14, wherein the clinical condition is selected from the group of asthma, nocturnal asthma, exercise-induced asthma, chronic obstructive pulmonary diseases (COPD), chronic and wheezy bronchitis, emphysema, , shortness of breath, respiratory tract infection and upper respiratory tract disease, rhinitis, allergic and seasonal rhinitis.
16. Method according to claim 15, which comprises a twice daily dosage regimen.
17. Method according to claim 15, which comprises a once daily dosage regimen.
18. Method according to claim 15, which comprises administration of a combination of the a pharmaceutical acceptable salt of glycopyrronium and ciclesonide in the same administration form

by inhalation from an inhaler and wherein each actuation provides a dose therapeutically effective for a twice daily dosing regimen or for a once daily dosing regimen.

19. Dry powder inhalation product comprising a pharmaceutical composition according to claim 13.